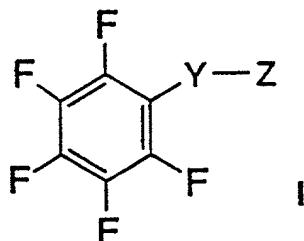


WHAT IS CLAIMED IS:

1. A pharmaceutical composition comprising a pharmaceutically acceptable excipient and a compound of formula I:

5



10

or a pharmaceutically acceptable salt thereof, wherein:

Y is -S(O)- or -S(O)₂;- and

Z is -NR¹R² or -OR³; wherein R¹ and R² are independently selected from

hydrogen,

substituted or unsubstituted (C1-C10)alkyl,

substituted or unsubstituted (C1-C10)alkoxy,

substituted or unsubstituted (C3-C6)alkenyl,

substituted or unsubstituted (C2-C6)heteroalkyl,

substituted or unsubstituted (C3-C6)heteroalkenyl,

substituted or unsubstituted (C3-C6)alkynyl,

substituted or unsubstituted (C3-C8)cycloalkyl,

substituted or unsubstituted (C5-C7)cycloalkenyl,

substituted or unsubstituted (C5-C7)cycloalkadienyl,

substituted or unsubstituted aryl,

substituted or unsubstituted aryloxy,

substituted or unsubstituted aryl-(C3-C8)cycloalkyl,

substituted or unsubstituted aryl-(C5-C7)cycloalkenyl,

substituted or unsubstituted aryloxy-(C3-C8)cycloalkyl,

substituted or unsubstituted aryl-(C1-C4)alkyl,

substituted or unsubstituted aryl-(C1-C4)alkoxy,

substituted or unsubstituted aryl-(C1-C4)heteroalkyl,

15

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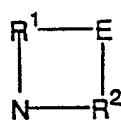
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substituted or unsubstituted aryl-(C3-C6)alkenyl,
 substituted or unsubstituted aryloxy-(C1-C4)alkyl,
 substituted or unsubstituted aryloxy-(C2-C4)heteroalkyl,
 substituted or unsubstituted heteroaryl,
 5 substituted or unsubstituted heteroaryloxy,
 substituted or unsubstituted heteroaryl-(C1-C4)alkyl,
 substituted or unsubstituted heteroaryl-(C1-C4)alkoxy,
 substituted or unsubstituted heteroaryl-(C1-C4)heteroalkyl,
 substituted or unsubstituted heteroaryl-(C3-C6)alkenyl,
 10 substituted or unsubstituted heteroaryloxy-(C1-C4)alkyl, and
 substituted or unsubstituted heteroaryloxy-(C2-C4)heteroalkyl,

wherein R¹ and R² may be connected by a linking group E to give a substituent of the formula



15 wherein E represents a bond, (C1-C4) alkylene, or (C1-C4) heteroalkylene, and the ring formed by R¹, E, R² and the nitrogen atom contains no more than 8 atoms; and where R³ is a substituted or unsubstituted aryl or heteroaryl group, wherein said compound I has pharmacological activity.

20 2. The composition of claim 1, wherein, in the compound of formula I,

Y is SO₂ and

Z is NR¹R²; wherein R² is optionally substituted aryl or optionally substituted heteroaryl.

25 3. The composition of claim 2, wherein R¹ is hydrogen or lower alkyl, R² is optionally substituted phenyl or optionally substituted pyridyl, and there is no linking group E between R¹ and R².

30 4. The composition of claim 3, wherein R¹ is hydrogen or methyl and R² is substituted phenyl, wherein the substituents on R², ranging in number from one to four, are

independently chosen from lower alkyl, hydroxy, lower alkoxy, amino optionally substituted with one or two lower alkyls, optionally substituted arylamino, optionally substituted heteroaryl amino, optionally substituted phenoxy, and halogen.

5 5. The composition of claim 4, wherein R¹ is hydrogen and R² is substituted phenyl, wherein the substituents on R² are independently chosen from amino, (lower)alkylamino, and di(lower)alkylamino, and are located at one or more of positions 3- and 4- of the phenyl ring, in relation to the sulfonamido group.

10 6. The composition of claim 5, wherein the compound is
4-(N,N-Dimethylamino)-1-pentafluorophenylsulfonamidobenzene,
4-(N,N-Diethylamino)-1-pentafluorophenylsulfonamidobenzene,
3-(N,N-Dimethylamino)-1-pentafluorophenylsulfonamidobenzene,
4-Amino-1-pentafluorophenylsulfonamidobenzene, or
15 4-(N,N-Dimethylamino)-1-pentafluorophenylsulfonamidobenzene hydrochloride.

7. The composition of claim 6, wherein the compound is
4-(N,N-Dimethylamino)-1-pentafluorophenylsulfonamidobenzene.

20 8. The composition of claim 2, wherein the compound is
Pentafluorophenylsulfonamidobenzene.

9. The composition of claim 3, wherein R¹ is hydrogen, and R² is phenyl substituted at positions 3- and 4-, in relation to the sulfonamido group, with a divalent moiety that forms a
25 5- or 6- membered ring together with carbons 3- and 4- of the phenyl ring.

10. The composition of claim 9, wherein the divalent moiety is: -OCH₂CH₂O-,
-OCH₂O-, -C=CNH-, or -C=NNH-.

30 11. The composition of claim 10, wherein the compound is
1,2-Ethylenedioxy-4-pentafluorophenylsulfonamidobenzene,

1,2-Methylenedioxy-4-pentafluorophenylsulfonamidobenzene,
5-Pentafluorophenylsulfonamidoindazole, or
5-Pentafluorophenylsulfonamidoindole.

5 12. The composition of claim 4, wherein R¹ is hydrogen, and the substituents on R² are independently selected from halogen, hydroxy, lower alkyl, lower alkoxy, amino, (lower)alkylamino, and di(lower)alkylamino.

10 13. The composition of claim 12, wherein the substituents on R² are independently selected from bromo, chloro, fluoro, hydroxy, methoxy, ethoxy, amino, and dimethylamino.

14. The composition of claim 13, wherein the substituents on R² are independently selected from bromo, chloro, fluoro, hydroxy, methoxy, and ethoxy.

15 15. The composition of claim 12, wherein the substituents on R² are at one or more of positions 3- and 4- of the phenyl ring, in relation to the sulfonamido group.

16. The composition of claim 15, wherein R² is monosubstituted phenyl.

20 17. The composition of claim 16, wherein the compound is
4-Methoxy-1-pentafluorophenylsulfonamidobenzene,
3-Hydroxy-1-pentafluorophenylsulfonamidobenzene,
4-Hydroxy-1-pentafluorophenylsulfonamidobenzene,
4-Ethoxy-1-pentafluorophenylsulfonamidobenzene,
25 3-Ethoxy-1-pentafluorophenylsulfonamidobenzene,
3-Phenoxy-1-pentafluorophenylsulfonamidobenzene,
3-Methoxy-1-pentafluorophenylsulfonamidobenzene, or
4-*tert*-Butoxy-1-pentafluorophenylsulfonamidobenzene.

30 18. The composition of claim 16, wherein the compound is
3-Chloro-1-pentafluorophenylsulfonamidobenzene, or

4-Chloro-1-pentafluoro-phenylsulfonamidobenzene.

19. The composition of claim 15, wherein R² is disubstituted phenyl.

5 20. The composition of claim 19, wherein the compound is
1,2-Dimethoxy-4-pentafluorophenylsulfonamidobenzene,
2-Hydroxy-1-methoxy-4-pentafluorophenylsulfonamidobenzene,
1,2-Dihydroxy-4-pentafluorophenylsulfonamidobenzene,
2-Hydroxy-1-methoxy-4-pentafluorophenylsulfonamidobenzene, monosodium salt, or
10 2-Hydroxy-1-methoxy-4-pentafluorophenylsulfonamidobenzene, monopotassium salt.

21. The composition of claim 19, wherein the compound is
2-Fluoro-1-methoxy-4-pentafluorophenylsulfonamidobenzene,
2-Bromo-1-methoxy-4-pentafluorophenylsulfonamidobenzene,
15 2-Chloro-1-methoxy-4-pentafluorophenylsulfonamidobenzene,
2-Fluoro-1-methoxy-4-pentafluorophenylsulfonamidobenzene, sodium salt, or
2-Fluoro-1-methoxy-4-pentafluorophenylsulfonamidobenzene, potassium salt.

22. The composition of claim 20, wherein the compound is
20 2-Hydroxy-1-methoxy-4-pentafluorophenylsulfonamidobenzene.

23. The composition of claim 20, wherein the compound is
2-Hydroxy-1-methoxy-4-pentafluorophenylsulfonamidobenzene, monosodium salt.

25 24. The composition of claim 21, wherein the compound is
2-Fluoro-1-methoxy-4-pentafluorophenylsulfonamidobenzene.

25. The composition of claim 21, wherein the compound is
2-Fluoro-1-methoxy-4-pentafluorophenylsulfonamidobenzene, sodium salt.

30 26. The composition of claim 21, wherein the compound is

2-Fluoro-1-methoxy-4-pentafluorophenylsulfonamidobenzene, potassium salt.

27. The composition of claim 12, wherein R² is a trisubstituted phenyl.

5 28. The composition of claim 21, wherein the compound is
2-Bromo-1-methoxy-4-pentafluorophenylsulfonamidobenzene, or
2-Chloro-1-methoxy-4-pentafluorophenylsulfonamidobenzene.

10 29. The composition of claim 12, wherein the compound is
1,2-Dimethyl-4-pentafluorophenylsulfonamidobenzene.

30. The composition of claim 1, wherein in the compound of formula I,

Y is SO₂ and

15 Z is NR¹R², where R¹ is hydrogen or lower alkyl, and R² is an unsubstituted or
optionally substituted naphthyl group.

31. The composition of claim 30, wherein the compound is
7-Hydroxy-2-pentafluorophenylsulfonamidonaphthalene.

20 32. The composition of claim 4, wherein R² is a phenyl group substituted by phenoxy or
optionally substituted phenoxy.

33. The composition of claim 32, wherein the compound is
3-Phenoxy-1-pentafluorophenylsulfonamidobenzene.

25

34. The composition of claim 3, wherein R² is a phenyl ring substituted by a heterocyclic
group at the 4- position, in relation to the sulfonamido group.

30 35. The composition of claim 17, wherein the compound is
4-Methoxy-1-pentafluorophenylsulfonamidobenzene.

36. The composition of claim 2, wherein R¹ and R² are covalently joined in a moiety that forms a 5- or 6- membered heterocyclic ring with the nitrogen atom of NR¹R².

37. The composition of claim 36, wherein R¹ is a -CH=CH- group linked to the 2-
5 position of the R² phenyl group, in relation to the sulfonamido group, forming an optionally substituted indole.

38. The composition of claim 37, wherein the compound is 1-pentafluorophenylsulfonylindole.

10

39. The composition of claim 36, wherein R¹ is a -(CH₂)₃- group linked to the 2- position of the R² phenyl group, in relation to the sulfonamido group, forming an optionally substituted 1,2,3,4-tetrahydroquinoline.

15 40. The composition of claim 39, wherein the compound is 1-pentafluorophenylsulfonyl-1,2,3,4-tetrahydroquinoline.

41. The composition of claim 2, wherein R¹ is an optionally substituted (C₂-C₁₀)alkyl or optionally substituted (C₂-C₆)heteroalkyl.

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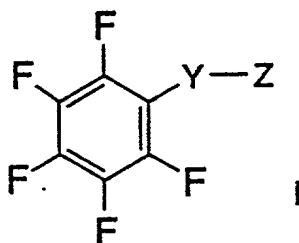
42. The composition of claim 41, wherein the compound is 2-Hydroxy-1-methoxy-4-[N-(5-hydroxypent-1-yl)pentafluorophenyl-sulfonamido]benzene, 4-Methoxy-1-[N-(2-propenyl)pentafluorophenylsulfonamido]benzene, 4-Methoxy-1-[N-(4-pentenyl)pentafluorophenylsulfonamido]benzene, 1-[N-(2,3-Dihydroxypropyl) pentafluorophenylsulfonamido]-4-methoxybenzene, 1-[N-(3,4-Dihydroxybutyl)pentafluorophenylsulfonamido]-4-methoxybenzene, 1-[N-(4,5-Dihydroxypentyl)pentafluorophenylsulfonamido]-4-methoxybenzene, 1-[N-(4-hydroxybutyl)pentafluorophenylsulfonamido]-4-methoxybenzene, or 4-Methoxy-1-[N-(5-hydroxypentyl)pentafluorophenylsulfonamido]benzene.

30

43. A method of treating or preventing a disease state characterized by abnormally high

levels of low density lipoprotein particles or cholesterol in the blood, which method comprises administering to a mammalian subject in need thereof a therapeutically effective amount of a composition containing a compound of formula I

5



10 or a pharmaceutically acceptable salt thereof, wherein:

Y is $-S(O)-$ or $-S(O)_2-$;

Z is $-NR^1R^2$ or $-OR^3$; where R^1 and R^2 are independently selected from hydrogen,

substituted or unsubstituted (C1-C10)alkyl,

substituted or unsubstituted (C1-C10)alkoxy,

substituted or unsubstituted (C3-C6)alkenyl,

substituted or unsubstituted (C2-C6)heteroalkyl,

substituted or unsubstituted (C3-C6)heteroalkenyl,

substituted or unsubstituted (C3-C6)alkynyl,

substituted or unsubstituted (C3-C8)cycloalkyl,

substituted or unsubstituted (C5-C7)cycloalkenyl,

substituted or unsubstituted (C5-C7)cycloalkadienyl,

substituted or unsubstituted aryl,

substituted or unsubstituted aryloxy,

20

substituted or unsubstituted aryl-(C3-C8)cycloalkyl,

substituted or unsubstituted aryl-(C5-C7)cycloalkenyl,

substituted or unsubstituted aryloxy-(C3-C8)cycloalkyl,

25

substituted or unsubstituted aryl-(C1-C4)alkyl,

substituted or unsubstituted aryl-(C1-C4)alkoxy,

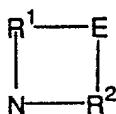
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substituted or unsubstituted aryl-(C1-C4)heteroalkyl,

substituted or unsubstituted aryl-(C3-C6)alkenyl,

substituted or unsubstituted aryloxy-(C1-C4)alkyl,
 substituted or unsubstituted aryloxy-(C2-C4)heteroalkyl,
 substituted or unsubstituted heteroaryl,
 substituted or unsubstituted heteroaryloxy,
 5 substituted or unsubstituted heteroaryl-(C1-C4)alkyl,
 substituted or unsubstituted heteroaryl-(C1-C4)alkoxy,
 substituted or unsubstituted heteroaryl-(C1-C4)heteroalkyl,
 substituted or unsubstituted heteroaryl-(C3-C6)alkenyl,
 substituted or unsubstituted heteroaryloxy-(C1-C4)alkyl, and
 10 substituted or unsubstituted heteroaryloxy-(C2-C4)heteroalkyl,

wherein R¹ and R² may be connected by a linking group E to give a substituent of the formula



15 wherein E represents a bond, (C1-C4) alkylene, or (C1-C4) heteroalkylene, and the ring formed by R¹, E, R² and the nitrogen contains no more than 8 atoms;
 and where R³ is optionally substituted aryl or optionally substituted heteroaryl.

44. The method of claim 43 wherein, in the compound of formula I,
 20 Y is SO₂ and
 Z is NR¹R²; where R² is optionally substituted aryl or optionally substituted heteroaryl.

45. The method of claim 44, wherein R¹ is hydrogen or lower alkyl, R² is optionally substituted phenyl, and there is no linking group E between R¹ and R².
 25

46. The method of claim 45, wherein R¹ is hydrogen or methyl, and the substituents on R² are independently chosen from lower alkyl, hydroxy, lower alkoxy, amino, amino optionally substituted with one or two lower alkyls, optionally substituted arylamino, 30 optionally substituted heteroaryl amino, optionally substituted phenoxy, and halogen.

47. The method of claim 46, wherein the compound is chosen from:

4-(N,N-Dimethylamino)-1-pentafluorophenylsulfonamidobenzene,
3-(N,N-Dimethylamino)-1-pentafluorophenylsulfonamidobenzene,
1,2-Ethylenedioxy-4-pentafluorophenylsulfonamidobenzene,
5 1,2-Methylenedioxy-4-pentafluorophenylsulfonamidobenzene,
1,2-Dimethoxy-4-pentafluorophenylsulfonamidobenzene,
2-Hydroxy-1-methoxy-4-pentafluorophenylsulfonamidobenzene,
2-Fluoro-1-methoxy-4-pentafluorophenylsulfonamidobenzene,
4-Methoxy-1-pentafluorophenylsulfonamidobenzene,
10 3-Hydroxy-1-pentafluorophenylsulfonamidobenzene,
4-Hydroxy-1-pentafluorophenylsulfonamidobenzene,
1,2-Dimethyl-4-pentafluorophenylsulfonamidobenzene,
4-(N,N-Diethylamino)-1-pentafluorophenylsulfonamidobenzene,
4-Amino-1-pentafluorophenylsulfonamidobenzene,
15 Pentafluorophenylsulfonamidobenzene,
5-Pentafluorophenylsulfonamidoindazole,
5-Pentafluorophenylsulfonamidoindole,
4-(N,N-Dimethylamino)-1-(N-methylpentafluorophenylsulfonamido)benzene,
4-(N,N-Dimethylamino)-1-(pentafluorophenylsulfonamido)benzene,
20 1,2-Dihydroxy-4-pentafluorophenylsulfonamidobenzene,
4-Ethoxy-1-pentafluorophenylsulfonamidobenzene,
3,5-Dimethoxy-1-pentafluorophenylsulfonamidobenzene,
3-Ethoxy-1-pentafluorophenylsulfonamidobenzene,
7-Hydroxy-2-pentafluorophenylsulfonamidonaphthalene,
25 3-Phenoxy-1-pentafluorophenylsulfonamidobenzene,
3-Methoxy-1-pentafluorophenylsulfonamidobenzene,
4(1-Morpholino)-1-pentafluorophenylsulfonamidobenzene,
5-Pentafluorophenylsulfonamido-1,2,3-trimethoxybenzene,
2-Hydroxy-1,3-methoxy-5-pentafluorophenylsulfonamidobenzene,
30 1,2-Dihydroxy-3-methoxy-5-pentafluorophenylsulfonamidobenzene,
5-Pentafluorophenylsulfonamido-1,2,3-trihydroxybenzene,

1,3-Dimethoxy-2-hydroxy-5-pentafluorophenylsulfonamidobenzene,
 1,2-Dihydroxy-3-methoxy-5-pentafluorophenylsulfonamidobenzene,
 5-Pentafluorophenylsulfonamido-1,2,3-trihydroxybenzene,
 3-Hydroxy-5-methoxy-1-pentafluorophenylsulfonamidobenzene,
 5 3,5-Dihydroxy-1-pentafluorophenylsulfonamidobenzene,
 2-Fluoro-1-methoxy-4-(N-methylpentafluorophenylsulfonamido)benzene,
 2-Bromo-1-methoxy-4-pentafluorophenylsulfonamidobenzene,
 2-Chloro-1-methoxy-4-pentafluorophenylsulfonamidobenzene,
 4-(N,N-Dimethylamino)-1-pentafluorophenylsulfonamidobenzene hydrochloride,
 10 3,4-Difluoro-1-pentafluorophenylsulfonamidobenzene,
 4-Trifluoromethoxy-1-pentafluorophenylsulfonamidobenzene,
 2-Chloro-5-pentafluorophenylsulfonamidopyridine,
 2-Hydroxy-1-methoxy-4-[N-(5-hydroxypentyl)pentafluorophenylsulfonamido]benzene,
 4-(1,1-Dimethyl)ethoxy-1-pentafluorophenylsulfonamidobenzene,
 15 2-Bromo-3-hydroxy-4-methoxy-1-pentafluorophenylsulfonamidobenzene,
 2-Bromo-4-methoxy-5-hydroxy-1-pentafluorophenylsulfonamidobenzene,
 1-Bromo-4-fluoro-5-methoxy-2-pentafluorophenylsulfonamidobenzene,
 2-Hydroxy-1-methoxy-4-pentafluorophenylsulfonamidobenzene; sodium salt,
 2-Hydroxy-1-methoxy-4-pentafluorophenylsulfonamidobenzene; potassium salt,
 20 2-Fluoro-1-methoxy-4-pentafluorophenylsulfonamidobenzene; sodium salt,
 2-Fluoro-1-methoxy-4-pentafluorophenylsulfonamidobenzene; potassium salt,
 3-Chloro-1-pentafluorophenylsulfonamidobenzene,
 4-Chloro-1-pentafluorophenylsulfonamidobenzene,
 3-Nitro-1-pentafluorophenylsulfonamidobenzene,
 25 4-Methoxy-1-pentafluorophenylsulfonamido-3-trifluoromethylbenzene,
 4-Methoxy-1-(N-(2-propenyl)pentafluorophenylsulfonamido)benzene,
 1-(N-(3-Butenyl)pentafluorophenylsulfonamido)-4-methoxybenzene,
 4-Methoxy-1-(N-(4-pentenyl)pentafluorophenylsulfonamido)benzene,
 1-(N-(2,3-Dihydroxypropyl)pentafluorophenylsulfonamido)-4-methoxybenzene,
 30 1-(N-(3,4-Dihydroxybutyl)pentafluorophenylsulfonamido)-4-methoxybenzene,
 1-(N-(4,5-Dihydroxypentyl)pentafluorophenylsulfonamido)-4-methoxybenzene,

1-(N-(4-hydroxybutyl)pentafluorophenylsulfonamido)-4-methoxybenzene,
 4-Methoxy-1-(N-(5-hydroxypentyl)pentafluorophenylsulfonamido)benzene,
 4-Methoxy-3-nitro-1-pentafluorophenylsulfonamidobenzene,
 3-Amino-4-methoxy-1-pentafluorophenylsulfonamidobenzene,
 5 4-Butoxy-1-pentafluorophenylsulfonamidobenzene,
 1-Pentafluorophenylsulfonamido-4-phenoxybenzene,
 4-Benzylxy-1-pentafluorophenylsulfonamidobenzene,
 4-Methylmercapto-1-pentafluorophenylsulfonamidobenzene,
 2-Methoxy-1-pentafluorophenylsulfonamidobenzene,
 10 4-Allyloxy-1-pentafluorophenylsulfonamidobenzene,
 1-Pentafluorophenylsulfonamido-4-propoxybenzene,
 4-(1-Methyl)ethoxy-1-pentafluorophenylsulfonamidobenzene,
 1-Pentafluorophenylsulfonyloxybenzene,
 1-Pentafluorophenylsulfonylindole,
 15 1-Pentafluorophenylsulfonyl-1,2,3,4-tetrahydroquinoline,
 2-Methoxy-5-pentafluorophenylsulfonamidopyridine,
 2-Fluoro-1-methoxy-4-pentafluorophenylsulfonamide,
 2-Anilino-3-pentafluorophenylsulfonamidopyridine.
 20

48. The method of claim 47, wherein the compound is chosen from:
 4-(N,N-Dimethylamino)-1-pentafluorophenylsulfonamidobenzene,
 3-(N,N-Dimethylamino)-1-pentafluorophenylsulfonamidobenzene,
 1,2-Ethylenedioxy-4-pentafluorophenylsulfonamidobenzene,
 25 2-Hydroxy-1-methoxy-4-pentafluorophenylsulfonamidobenzene,
 2-Fluoro-1-methoxy-4-pentafluorophenylsulfonamidobenzene,
 4-Methoxy-1-pentafluorophenylsulfonamidobenzene,
 3-Hydroxy-1-pentafluorophenylsulfonamidobenzene,
 4-Hydroxy-1-pentafluorosulfonamidobenzene,
 30 1,2-Dimethyl-4-pentafluorophenylsulfonamidobenzene,
 5-Pentafluorophenylsulfonamidoindole,

4-(N,N-Dimethylamino)-1-(N-methylpentafluorophenylsulfonamido)benzene,
4-Ethoxy-1-pentafluorophenylsulfonamidobenzene,
3-Methoxy-1-pentafluorophenylsulfonamidobenzene,
2-Bromo-1-methoxy-4-pentafluorophenylsulfonamidobenzene,
5 2-Chloro-1-methoxy-4-pentafluorophenylsulfonamidobenzene,
2-Bromo-3-hydroxy-4-methoxy-1-pentafluorophenylsulfonamidobenzene,
2-Bromo-4-methoxy-5-hydroxy-1-pentafluorophenylsulfonamidobenzene,
1-Bromo-4-fluoro-5-methoxy-2-pentafluorophenylsulfonamidobenzene,
2-Hydroxy-1-methoxy-4-pentafluorophenylsulfonamidobenzene; monosodium salt,
10 2-Hydroxy-1-methoxy-4-pentafluorophenylsulfonamidobenzene; monopotassium salt,
2-Fluoro-1-methoxy-4-pentafluorophenylsulfonamidobenzene; sodium salt,
2-Fluoro-1-methoxy-4-pentafluorophenylsulfonamidobenzene; potassium salt,
4-Chloro-1-pentafluorophenylsulfonamidobenzene, and
3-Amino-4-methoxy-1-pentafluorophenylsulfonamidobenzene.

15 49. The method of claim 48, wherein the compound is:

2-Hydroxy-1-methoxy-4-pentafluorophenylsulfonamidobenzene,
2-Fluoro-1-methoxy-4-pentafluorophenylsulfonamidobenzene,
4-Methoxy-1-pentafluorophenylsulfonamidobenzene,
20 1,2-Dimethyl-4-pentafluorophenylsulfonamidobenzene,
5-Pentafluorophenylsulfonamidoindole,
4-Ethoxy-1-pentafluorophenylsulfonamidobenzene,
2-Bromo-1-methoxy-4-pentafluorophenylsulfonamidobenzene,
2-Chloro-1-methoxy-4-pentafluorophenylsulfonamidobenzene,
25 2-Bromo-3-hydroxy-4-methoxy-1-pentafluorophenylsulfonamidobenzene,
2-Bromo-4-methoxy-5-hydroxy-1-pentafluorophenylsulfonamidobenzene,
1-Bromo-4-fluoro-5-methoxy-2-pentafluorophenylsulfonamidobenzene,
2-Hydroxy-1-methoxy-4-pentafluorophenylsulfonamidobenzene; monosodium salt,
2-Hydroxy-1-methoxy-4-pentafluorophenylsulfonamidobenzene; monopotassium salt,
30 2-Fluoro-1-methoxy-4-pentafluorophenylsulfonamidobenzene; sodium salt,
2-Fluoro-1-methoxy-4-pentafluorophenylsulfonamidobenzene; potassium salt, or

3-Amino-4-methoxy-1-pentafluorophenylsulfonamidobenzene.

50. The method of claim 49, wherein the compound is
2-Hydroxy-1-methoxy-4-pentafluorophenylsulfonamidobenzene,
5 2-Hydroxy-1-methoxy-4-pentafluorophenylsulfonamidobenzene; monosodium salt, or
2-Hydroxy-1-methoxy-4-pentafluorophenylsulfonamidobenzene; monopotassium salt.

51. The method of claim 49, wherein the compound is
2-Fluoro-1-methoxy-4-pentafluorophenylsulfonamidobenzene.

10 52. The method of claim 49, wherein the compound is
2-Fluoro-1-methoxy-4-pentafluorophenylsulfonamidobenzene, sodium salt.

53. The method of claim 49, wherein the compound is
15 2-Fluoro-1-methoxy-4-pentafluorophenylsulfonamidobenzene, potassium salt.

54. The method of claim 43 wherein the disease state is atherosclerosis.

55. The method of claim 43 wherein the disease state is pancreatitis.

20 56. The method of claim 43 wherein the disease state is hypercholesterolemia.

57. The method of claim 43 wherein the disease state is hyperlipoproteinemia.

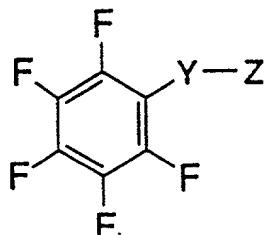
25 58. The method of claim 43 wherein the composition is administered orally.

59. The method of claim 43 wherein the subject is human.

60. The method of claim 43 wherein the composition is administered in combination with
30 a therapeutically effective amount of a hypolipemic agent or a hypocholesterolemic agent that
is not represented by formula I.

61. A compound having the formula I:

5



or a pharmaceutically acceptable salt thereof, wherein:

10 Y is -S(O)- or -S(O₂)-; and

Z is NR¹R², wherein R² is an optionally substituted aryl or heteroaryl group, and R¹ is selected from

hydrogen,

substituted or unsubstituted (C1-C10)alkyl,

substituted or unsubstituted (C1-C10)alkoxy,

substituted or unsubstituted (C3-C6)alkenyl,

substituted or unsubstituted (C2-C6)heteroalkyl,

substituted or unsubstituted (C3-C6)heteroalkenyl,

substituted or unsubstituted (C3-C6)alkynyl,

substituted or unsubstituted (C3-C8)cycloalkyl,

substituted or unsubstituted (C5-C7)cycloalkenyl,

substituted or unsubstituted (C5-C7)cycloalkadienyl,

substituted or unsubstituted aryl,

substituted or unsubstituted aryloxy,

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substituted or unsubstituted aryl-(C3-C8)cycloalkyl,

substituted or unsubstituted aryl-(C5-C7)cycloalkenyl,

substituted or unsubstituted aryloxy-(C3-C8)cycloalkyl,

substituted or unsubstituted aryl-(C1-C4)alkyl,

substituted or unsubstituted aryl-(C1-C4)alkoxy,

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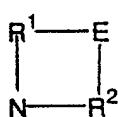
substituted or unsubstituted aryl-(C1-C4)heteroalkyl,

substituted or unsubstituted aryl-(C3-C6)alkenyl,

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substituted or unsubstituted aryloxy-(C1-C4)alkyl,
 substituted or unsubstituted aryloxy-(C2-C4)heteroalkyl,
 substituted or unsubstituted heteroaryl,
 substituted or unsubstituted heteroaryloxy,
 5 substituted or unsubstituted heteroaryl-(C1-C4)alkyl,
 substituted or unsubstituted heteroaryl-(C1-C4)alkoxy,
 substituted or unsubstituted heteroaryl-(C1-C4)heteroalkyl,
 substituted or unsubstituted heteroaryl-(C3-C6)alkenyl,
 substituted or unsubstituted heteroaryloxy-(C1-C4)alkyl, and
 10 substituted or unsubstituted heteroaryloxy-(C2-C4)heteroalkyl,

wherein R¹ and R² may be connected by a linking group E to give a substituent of the formula



15 wherein E represents a bond, (C1-C4) alkylene, or (C1-C4) heteroalkylene, and the ring formed by R¹, E, R² and the nitrogen contains no more than 8 atoms;

provided that:

in the case that Y is -S(O₂)-, and R¹ is hydrogen or methyl, then R² is substituted phenyl or heteroaryl group;

20 in the case that Y is -S(O₂)- and R² is a ring system chosen from 1-naphthyl, 5-quinolyl, or 4-pyridyl, then either R¹ is not hydrogen or R² is substituted by at least one substituent that is not hydrogen;

25 in the case that Y is -S(O₂)-, R² is phenyl, and R¹ is a propylene unit attaching the nitrogen of -NR¹R²- to the 2- position of the phenyl ring in relation to the sulfonamido group to form a 1,2,3,4-tetrahydroquinoline system, and one or more of the remaining valences on the bicyclic system so formed is substituted with at least one substituent that is not hydrogen;

30 in the case that Y is -S(O₂)- and R² is phenyl substituted with 3-(1-hydroxyethyl), 3-dimethylamino, 4-dimethylamino, 4-phenyl, 3-hydroxy, 3-hydroxy-4-diethylaminomethyl, 3,4-methylenedioxy, 3,4-ethylenedioxy, 2-(1-pyrrolyl), or 2-methoxy-4-(1-morpholino), then either R¹ is not hydrogen or when R¹ is hydrogen, one or more of the remaining valences

on the phenyl ring of R² is substituted with a substituent that is not hydrogen;

in the case that Y is -S(O₂)- and R² is 2-methylbenzothiazol-5-yl, 6-hydroxy-4-methyl-pyrimidin-2-yl, 3-carbomethoxy pyrazin-2-yl, 5-carbomethoxypyrazin-2-yl, 4-carboethoxy-1-phenylpyrazol-5-yl, 3-methylpyrazol-5-yl, 5 4-chloro-2-methylthiopyrimidin-6-yl, 2-trifluoromethyl-1,3,4-thiadiazol-5-yl, 5,6,7,8-tetrahydro-2-naphthyl, 4-methylthiazol-2-yl, 6,7-dihydroindan-5-yl, 7-chloro-5-methyl-1,8-naphthyridin-2-yl, 5,7-dimethyl-1,8-naphthyridin-2-yl, or 3-cyanopyrazol-4-yl, then R¹ is a group other than hydrogen;
wherein said compound has pharmacological activity.

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62. The compound of claim 61, wherein R¹ is hydrogen or lower alkyl, Y is -S(O₂)-, and there is no linking group E between R¹ and R².

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63. The compound of claim 62, wherein R¹ is hydrogen or methyl and R² is substituted phenyl, wherein the substituents on R², ranging in number from one to four, are independently chosen from lower alkyl, hydroxy, lower alkoxy, amino optionally substituted with one or two lower alkyls, optionally substituted arylamino, optionally substituted heteroaryl amino, optionally substituted phenoxy, and halogen.

64. The compound of claim 63, wherein R¹ is hydrogen and R² is substituted phenyl, wherein the substituents on R² are independently chosen from amino, (lower)alkylamino, and di(lower)alkylamino, and are located at one or more of positions 3- and 4- of the phenyl ring, in relation to the sulfonamido group.

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65. The compound of claim 64, wherein the compound is 4-(N,N-Diethylamino)-1-pentafluorophenylsulfonamidobenzene, or 4-Amino-1-pentafluorophenylsulfonamidobenzene.

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66. The compound of claim 62, wherein R¹ is hydrogen, and R² is phenyl substituted at positions 3- and 4-, in relation to the sulfonamido group, with a divalent moiety that forms a 5- or 6- membered ring together with carbons 3- and 4- of the phenyl ring.

67. The compound of claim 66, wherein the divalent moiety is: -C=CNH-, or -C=NNH-.

68. The compound of claim 67, wherein the compound is 5-
5 Pentafluorophenylsulfonamidoindazole or 5-Pentafluorophenylsulfonamidoindole.

69. The compound of claim 63, wherein R¹ is hydrogen, and the substituents on R² are independently selected from halogen, hydroxy, lower alkyl, lower alkoxy, amino, (lower)alkylamino, and di(lower)alkylamino.

10 70. The compound of claim 69, wherein the substituents on R² are independently selected from bromo, chloro, fluoro, hydroxy, methoxy, ethoxy, amino, or dimethylamino.

15 71. The compound of claim 70, wherein the substituents on R² are independently selected from bromo, chloro, fluoro, hydroxy, methoxy, and ethoxy.

72. The compound of claim 71, wherein the substituents on R² are at one or more of positions 3- and 4- of the phenyl ring, in relation to the sulfonamido group.

20 73. The compound of claim 72, wherein R² is monosubstituted phenyl.

74. The compound of claim 73, wherein the compound is
4-Methoxy-1-pentafluorophenylsulfonamidobenzene,
3-Hydroxy-1-pentafluorophenylsulfonamidobenzene,
25 4-Hydroxy-1-pentafluorophenylsulfonamidobenzene,
4-Ethoxy-1-pentafluorophenylsulfonamidobenzene,
3-Ethoxy-1-pentafluorophenylsulfonamidobenzene, or
3-Methoxy-1-pentafluorophenylsulfonamidobenzene.

30 75. The compound of claim 73, wherein the compound is
3-Chloro-1-pentafluorophenylsulfonamidobenzene, or

4-Chloro-1-pentafluorophenylsulfonamidobenzene.

76. The compound of claim 72, wherein R² is disubstituted phenyl.

5 77. The compound of claim 76, wherein the compound is

1,2-Dimethoxy-4-pentafluorophenylsulfonamidobenzene,

2-Hydroxy-1-methoxy-4-pentafluorophenylsulfonamidobenzene,

1,2-Dihydroxy-4-pentafluorophenylsulfonamidobenzene,

2-Hydroxy-1-methoxy-4-pentafluorophenylsulfonamidobenzene, monosodium salt, or

10 2-Hydroxy-1-methoxy-4-pentafluorophenylsulfonamidobenzene, monopotassium salt.

78. The compound of claim 76, wherein the compound is

2-Fluoro-1-methoxy-4-pentafluorophenylsulfonamidobenzene,

2-Bromo-1-methoxy-4-pentafluorophenylsulfonamidobenzene,

15 2-Chloro-1-methoxy-4-pentafluorophenylsulfonamidobenzene,

2-Fluoro-1-methoxy-4-pentafluorophenylsulfonamidobenzene, sodium salt, or

2-Fluoro-1-methoxy-4-pentafluorophenylsulfonamidobenzene, potassium salt.

79. The compound of claim 77, wherein the compound is

20 2-Hydroxy-1-methoxy-4-pentafluorophenylsulfonamidobenzene, or

2-Hydroxy-1-methoxy-4-pentafluorophenylsulfonamidobenzene, monosodium salt.

80. The compound of claim 78, wherein the compound is

2-Fluoro-1-methoxy-4-pentafluorophenylsulfonamidobenzene, or

25 2-Fluoro-1-methoxy-4-pentafluorophenylsulfonamidobenzene, sodium salt.

81. The compound of claim 80, wherein the compound is

2-Fluoro-1-methoxy-4-pentafluorophenylsulfonamidobenzene.

30 82. The compound of claim 80, wherein the compound is

2-Fluoro-1-methoxy-4-pentafluorophenylsulfonamidobenzene, sodium salt.

83. The compound of claim 72, wherein R² is trisubstituted phenyl.

84. The compound of claim 78, wherein the compound is
5 2-Bromo-1-methoxy-4-pentafluorophenylsulfonamidobenzene, or
2-Chloro-1-methoxy-4-pentafluorophenylsulfonamidobenzene.

85. The compound of claim 69, wherein the compound is
1,2-Dimethyl-4-pentafluorophenylsulfonamidobenzene.

10 86. The compound of claim 63, wherein the compound is
3-Phenoxy-1-pentafluorophenylsulfonamidobenzene.

15 87. The compound of claim 62, wherein R² is a phenyl ring substituted by a heterocyclic
group at the 4- position, in relation to the sulfonamido group.

88. The compound of claim 74, wherein the compound is
4-Methoxy-1-pentafluorophenylsulfonamidobenzene.

20 89. The compound of claim 61, wherein R¹ and R² are covalently joined in a moiety that
forms a 5- or 6- membered heterocyclic ring with the nitrogen atom of NR¹R².

90. The compound of claim 89, wherein R¹ is a -CH=CH- group linked to the 2- position
of the R² phenyl group, in relation to the sulfonamido group, forming an optionally
25 substituted indole.

91. The compound of claim 90, wherein the compound is
1-(Pentafluorophenylsulfonyl)indole.

30 92. The compound of claim 89, wherein R¹ is a -(CH₂)₃- group linked to the 2- position
of the R² phenyl group, in relation to the sulfonamido group, forming an optionally

substituted 1,2,3,4-tetrahydroquinoline.

93. The compound of claim 61, wherein the compound is

2-Hydroxy-1-methoxy-4-[N-(5-hydroxypent-1-yl)pentafluorophenylsulfonamido]benzene,

5 4-Methoxy-1-[N-(2-propenyl)pentafluorophenylsulfonamido]benzene,

4-Methoxy-1-[N-(4-pentenyl)pentafluorophenylsulfonamido]benzene,

1-[N-(2,3-Dihydroxypropyl) pentafluorophenylsulfonamido]-4-methoxybenzene,

1-[N-(3,4-Dihydroxybutyl)pentafluorophenylsulfonamido]-4-methoxybenzene,

1-[N-(4,5-Dihydroxypentyl)pentafluorophenylsulfonamido]-4-methoxybenzene,

10 1-[N-(4-hydroxybutyl)pentafluorophenylsulfonamido]-4-methoxybenzene, or

4-Methoxy-1-[N-(5-hydroxypentyl)pentafluorophenylsulfonamido]benzene.

94. The composition of any of claim 1-42, or a method of any of claim 43-60, or a

compound of any of claim 61-93, wherein the compound regulates LDL receptor gene

expression.